Amendments To The Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

In the Claims:

What is claimed is:

1. (Original) A compound of formula (I):

$$(R^4)_r$$
 $(R^2)_n$
 $(R^2)_n$
 $(R^2)_m$
 $(R^3)_m$

wherein:

 R^1 represents hydrogen, $-C_{1-6}$ alkyl, $-C_{1-6}$ alkoxy, $-C_{3-8}$ cycloalkyl, $-C_{1-6}$ alkyl- C_{3-8} cycloalkyl, aryl, heterocyclyl, heteroaryl, $-C_{1-6}$ alkyl-aryl, $-C_{1-6}$ alkyl-heteroaryl, $-C_{1-6}$ alkyl-heterocyclyl, -aryl-aryl, -aryl-heteroaryl, -aryl-heterocyclyl, heteroaryl-aryl, -heterocyclyl-aryl, -heterocyclyl-heterocyclyl, heterocyclyl-heterocyclyl,

wherein R¹ may be optionally substituted by one or more substituents which may be the same or different, and which are selected from the group consisting of halogen, hydroxy, COOR¹⁵, cyano, -C₁₋₆ alkyl-cyano, nitro, oxo, trifluoromethyl, trifluoromethoxy, fluoromethoxy, difluoromethoxy, C₁₋₆ alkyl (optionally substituted by a COOR¹⁵ group), C₂₋₆ alkenyl (optionally substituted by a COOR¹⁵ group), C₂₋₆ alkynyl (optionally substituted by a COOR¹⁵ group), C₁₋₆ alkoxy (optionally substituted by a COOR¹⁵ group), pentafluoroethyl, C₁₋₆ alkoxy, C₂₋₆ alkenoxy, aryl, arylC₁₋₆ alkyl, -CO-aryl (optionally substituted by a halogen atom), -CO-heteroaryl, -C₁₋₆ alkyl-CO-aryl, arylC₁₋₆ alkoxy, C₁₋₆ alkylthio, C₁₋₆ alkoxyC₁₋₆ alkyl, C₃₋₇ cycloalkyl, C₃₋₇ cycloalkylC₁₋₆ alkoxy, C₁₋₈ 6 alkoxycarbonyl, C₁₋₆ alkylsulfonyl, C₁₋₆ alkylsulfinyl, C₁₋₆ alkylsulfonyloxy, C₁₋₆ alkylsulfonylC₁₋₆ alkyl, sulfonyl, arylsulfonyl, arylsulfonyloxy, arylsulfonylC₁₋₆ alkyl, aryloxy, C₁₋₆ alkylsulfonamido, C₁₋₆ alkylamido, C₁₋₆ alkylsulfonamidoC₁₋₆ alkyl, C₁₋₆ alkylamidoC₁₋₆ alkyl, arylsulfonamido, arylcarboxamido, arylsulfonamidoC₁₋₆ alkyl, arylcarboxamidoC₁₋₆ alkyl, aroyl, aroylC₁₋₆ alkyl, arylC₁₋₆ alkanoyl, or a group -COR¹⁵, -NR¹⁵R¹⁶, -CONR¹⁵R¹⁶, -NR¹⁵COR¹⁶, -NR¹⁵SO₂R¹⁶ or -SO₂NR¹⁵R¹⁶, wherein R¹⁵ and R¹⁶ independently represent hydrogen, C₁₋₆ alkyl or C₃₋₈ cycloalkyl or together may be fused to form a 5- to 7- membered non-aromatic heterocyclic ring optionally interrupted by an O or S atom and optionally substituted by a halogen, C_{1.6} alkyl or -C_{1.} 6 alkylC₁₋₆ alkoxy group;

Z represents a bond, CO, -CON(R¹⁰)- or SO₂, such that when R¹ represents hydrogen, Z represents CONR¹⁰;

p is 1 or 2;

m, n and r independently represent 0, 1 or 2;

 R^2 represents halogen, C_{1-6} alkyl, C_{1-6} alkoxy, cyano, amino or trifluoromethyl, such that when n represents 2, two R^2 groups may instead be linked to form a phenyl ring; R^4 represents C_{1-6} alkyl, such that when r represents 2, two R^4 groups may instead be linked to form a CH_2 , $(CH_2)_2$ or $(CH_2)_3$ group;

R¹⁰ represents hydrogen or C₁₋₆ alkyl, or R¹⁰, together with R¹ forms a heterocyclic group;

R³ represents -(CH₂)_q-NR¹¹R¹² or a group of formula (i):

$$--(CH_2)_f$$
 $(R^{14})_k$ (i)

wherein q is 2, 3 or 4;

R¹¹ and R¹² independently represent C₁₋₆ alkyl or C₃₋₈ cycloalkyl or together with the nitrogen atom to which they are attached represent an N-linked nitrogen containing heterocyclyl group optionally substituted by one or more R¹⁷ groups;

 R^{13} represents hydrogen, C_{1-6} alkyl, $-C_{1-6}$ alkyl- C_{1-6} alkoxy, C_{3-8} cycloalkyl, $-C_{1-6}$ alkyl-aryl or heterocyclyl;

 R^{14} and R^{17} independently represent halogen, C_{1-6} alkyl, haloalkyl, OH, di C_{1-6} alkylamino, C_{1-6} alkoxy or heterocyclyl;

f and k independently represent 0, 1 or 2;

g is 0, 1 or 2 and h is 0, 1, 2 or 3, such that g and h cannot both be 0; with the proviso that when m represents 1, n and r both represent 0 and R^3 represents $-(CH_2)_3-N$ -piperidine or $-(CH_2)_3-N$ (ethyl)₂, R^1-Z represents a group other than methyl, - $CO-O-C(CH_3)_3$ or benzyl;

and with the proviso that when m, n and r all represent 0, p represents 1, R^3 represents $-(CH_2)_3$ -N-pyrrolidine or $-(CH_2)_3$ -N-piperidine, R^1 represents benzyl, Z represents a group other than a bond;

and with the proviso that when m, n and r all represent 0, p represents 1, R^3 represents— $(CH_2)_3$ -N-piperidine, R^1 represents isopropyl, Z represents a group other than a bond;

and with the proviso that when m represents 1, n and r both represent 0, p represents 1, R^3 represents— $(CH_2)_3$ -N-piperidine, R^1 represents methyl, isopropyl, aryl or benzyl, Z represents a group other than a bond;

and with the proviso that when m and n both represent 0, R^3 represents $-(CH_2)_3$ - $N(ethyl)_2$, p represents 1, r represents 2 and R^1 and R^4 both represent methyl, Z represents a group other than a bond;

or a pharmaceutically acceptable salt thereof.

2. (Original) A compound according to claim 1 which is a compound of formula E1-E503 or a pharmaceutically acceptable salt thereof.

3. (Currently Amended) A pharmaceutical composition which comprises the compound of formula (I) as defined in claim 1 or claim 2 or a pharmaceutically acceptable salt thereof and a pharmaceutically acceptable carrier or excipient.

4. - 6. (Cancelled)

- 7. (Currently Amended) A method of treatment of neurological diseases <u>or inflammatory diseases</u> of the upper respiratory tract which comprises administering to a host in need thereof an effective amount of a compound of formula (I) as defined in claim 1 or claim 2 or a pharmaceutically acceptable salt thereof.
- 8. (Currently Amended) A pharmaceutical composition for use in the treatment of neurological diseases or inflammatory diseases of the upper respiratory tract which comprises the compound of formula (I) as defined in claim 1 or claim 2 or a pharmaceutically acceptable salt thereof and a pharmaceutically acceptable carrier.
- 9. (New) A pharmaceutical composition which comprises the compound of claim 2 or a pharmaceutically acceptable salt thereof and a pharmaceutically acceptable carrier or excipient.
- 10. (New) A pharmaceutical composition for the treatment of neurological diseases or inflammatory diseases of the upper respiratory tract which comprises the compound of claim 2 or a pharmaceutically acceptable salt thereof and a pharmaceutically acceptable carrier.
- 11. (New) A method of treatment of neurological diseases or inflammatory diseases of the upper respiratory tract which comprises administering to a host in need thereof an effective amount of a compound of claim 2 or a pharmaceutically acceptable salt thereof.